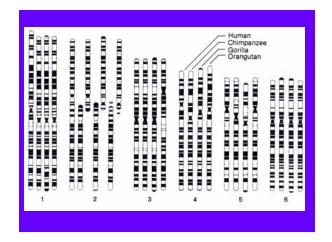
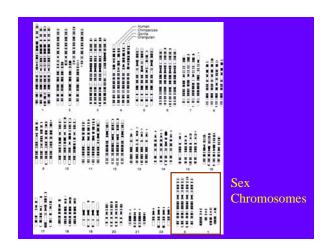


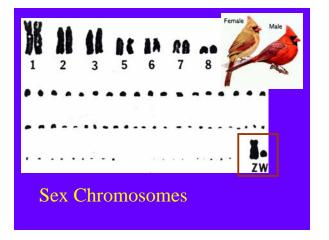
CHROMOSOMAL VARIABILITY

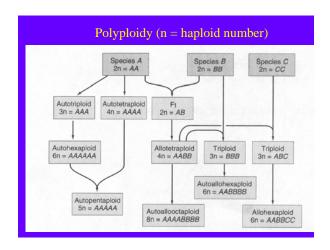
Ignored, but important for conservation:

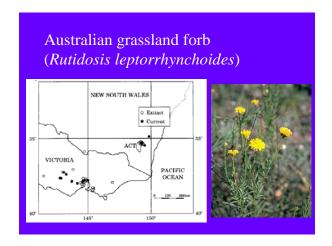
- (1) Associated with reduced fertility.
- (2) Taxa more likely to be threatened are also more likely to have more chromosomal variability.
 - (a) Small population size
 - (b) Complex social structure

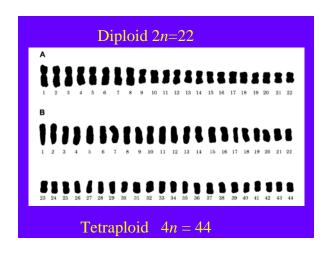


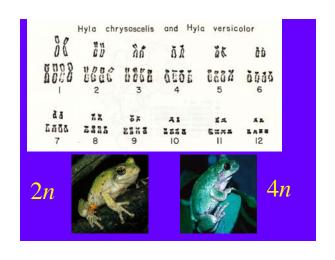


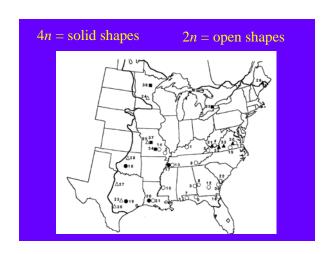


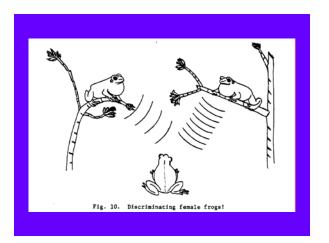


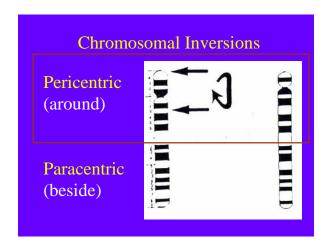




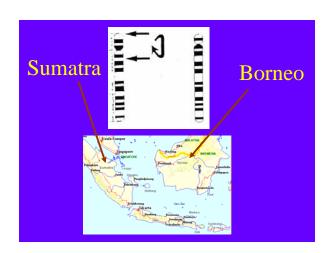












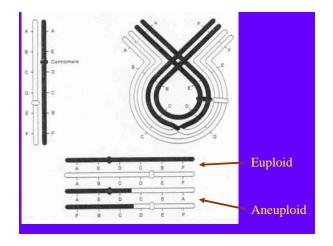
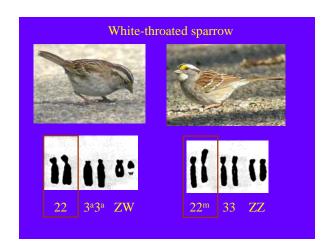
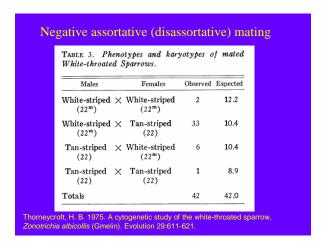
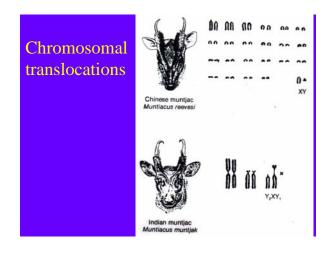


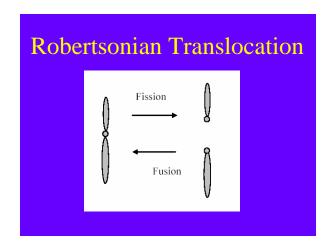
Table 3.2. Chromosomal inversion polymorphisms in the orangutan (Ryder and Chemnick 1993). The inversion in chromosome-2 distinguishes the Sumatran (S) and Bornean (B) subspecies. The two inversion types in chromosome-9 (C and R) are polymorphic in both subspecies.

	Chromosome 2		Chromosome 9			
	BB	SB	SS	CC	CR	RR
Wild born	51	0	41	67	22	3
Zoo born	90	44	82	71	34	3









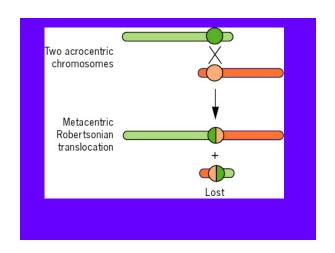
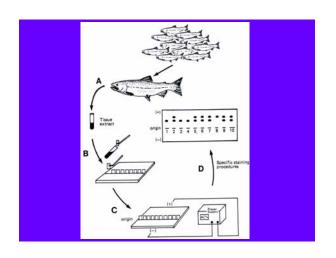


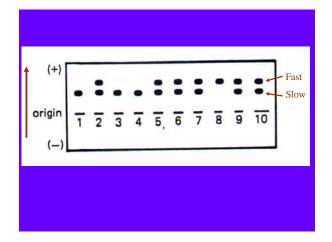
Table 3.3. Litter sizes produced by mice heterozygous for Robertsonian translocations characteristic of three different chromosomal races (AA, POS, and UV). From Hauffe and Searle (1998).

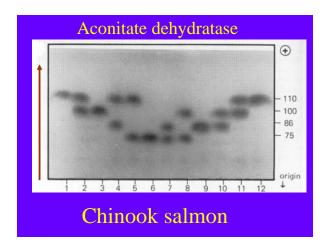
Ξ	Female	Male	No. litters	Litter size
	AA	AA (control)	17	6.7 ± 0.8
	AA	(AA x POS)	16	4.1 ± 0.4
	AA	(AA x UV)	18	2.6 ± 0.3
	AA	(UV x POS)	19	$\textbf{3.8} \pm \textbf{0.3}$
1	AA (control)	AA	18	6.8 ± 0.4
((AA x POS)	AA	7	1.0 ± 0
	(AA x UV)	AA	10	$\textbf{3.1} \pm \textbf{0.6}$
-	(POS x UV)	AA	11	4.0 ± 0.5

Protein Electrophoresis (Allozymes)

Time Period	Primary techniques
1900-1970	Laboratory matings and chromosomes
1970s	Protein electrophoresis (allozymes)
1980s	Mitochondrial DNA
1990s	Nuclear DNA
2000s	Genomics

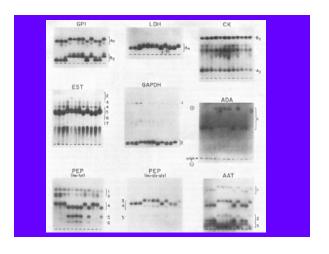


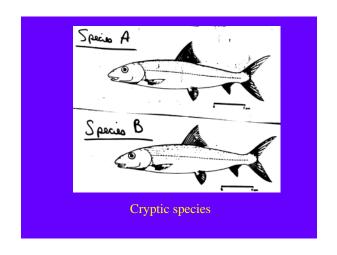


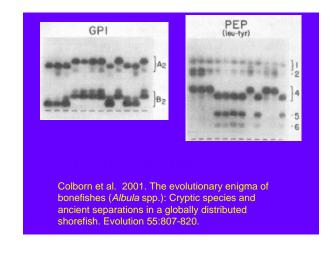


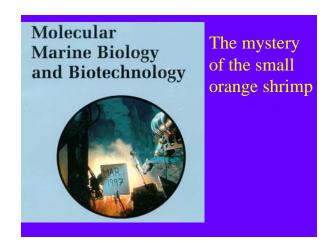
Are allozymes obsolete? Imagine for sake of argument that DNA sequencing methods had been widely employed for the past thirty years, and that only recently had protein electophoretic approaches been introduced. No doubt a headlong rush into allozyme techniques would ensue, on justifiable grounds that: (a) the methods are cost effective and technically simple; (b) the molecular variants represent independent Mendelian polymorphisms at numerous loci scattered around the genome (rather than tightly linked variants in a single sequenced region of DNA); and (c) the amino acid replacement substitutions revealed in the protein assays might bring molecular evolutionists closer to the real "stuff" of adaptive evolution. John Avise (1994)

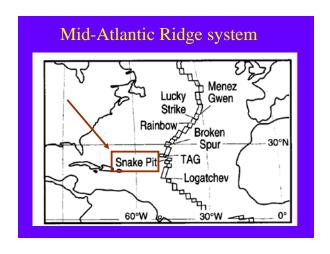
















Allleave	Genotype frequencies			
Allozyme locus and shrimp form	C/C	, C/R	R/R	
Pgm				
"Small orange"	24 (22.5)	4 (7.1)	1 (0.2)	
R. exoculata	23 (22.5)	6 (6.9)	1 (0.4)	
Pgi				
"Small orange"	9 (8.4)	14 (15.2)	7 (6.4)	
R. exoculata	6 (6.4)	16 (15.2)	8 (8.4)	
Gota				
"Small orange"	22 (22.5)	8 (7.1)	0 (0.4)	
R. exoculata	25 (25.2)	5 (4.6)	0 (0.2)	
Ap				
"Small orange"	27 (27.1)	3 (2.8)	0 (0.1)	
R. exoculata	24 (22.5)	5 (7.1)	1 (0.4)	

Small orange shrimp are juvenile *Rimicaris exoculata!*

9. Black and Johnson (1979) reported an highly unusual pattern of inheritance of allozyme polymorphisms in the intertidal anemone Actina tenebrosa from Rottnest Island in Western Australia. This species is viviparous, and up to 5 young are brooded by adults at a time until they are released as relatively large juveniles. The following parental and progeny genotypes were found at three allozyme loci:

Intertidal anemone



Asexual reproduction

	Parental	No. of	Progeny genotypes		
Locus	genotype	broods	FF	FS	SS
MDH	FF	25	68	0	0
	FS	53	0	158	0
	SS	11	0	0	35
PGM	FF	44	145	0	0
	FS	9	0	33	0
SOD	FF	71	225	0	0
	FS	18	0	50	0
	SS	1	0	0	2